- 1 AN ACT relating to drug paraphernalia.
- 2 Be it enacted by the General Assembly of the Commonwealth of Kentucky:
- 3 → Section 1. KRS 218A.010 is amended to read as follows:
- 4 As used in this chapter:
- 5 (1) "Administer" means the direct application of a controlled substance, whether by
- 6 injection, inhalation, ingestion, or any other means, to the body of a patient or
- 7 research subject by:
- 8 (a) A practitioner or by his or her authorized agent under his or her immediate
- 9 supervision and pursuant to his or her order; or
- 10 (b) The patient or research subject at the direction and in the presence of the
- 11 practitioner;
- 12 (2) "Anabolic steroid" means any drug or hormonal substance chemically and
- pharmacologically related to testosterone that promotes muscle growth and includes
- those substances classified as Schedule III controlled substances pursuant to KRS
- 15 218A.020 but does not include estrogens, progestins, and anticosteroids;
- 16 (3) "Cabinet" means the Cabinet for Health and Family Services;
- 17 (4) "Carfentanil" means any substance containing any quantity of carfentanil, or any of
- its salts, isomers, or salts of isomers;
- 19 (5) "Certified community based palliative care program" means a palliative care
- 20 program which has received certification from the Joint Commission;
- 21 (6) "Child" means any person under the age of majority as specified in KRS 2.015;
- 22 (7) "Cocaine" means a substance containing any quantity of cocaine, its salts, optical
- and geometric isomers, and salts of isomers;
- 24 (8) "Controlled substance" means methamphetamine, or a drug, substance, or
- 25 immediate precursor in Schedules I through V and includes a controlled substance
- analogue;
- 27 (9) (a) "Controlled substance analogue," except as provided in paragraph (b) of this

subsection, means a substance:

1

27

2	1.	The chemical structure of which is substantially similar to the structure
3		of a controlled substance in Schedule I or II; and
4	2.	Which has a stimulant, depressant, or hallucinogenic effect on the
5		central nervous system that is substantially similar to or greater than the
6		stimulant, depressant, or hallucinogenic effect on the central nervous
7		system of a controlled substance in Schedule I or II; or
8	3.	With respect to a particular person, which such person represents or
9		intends to have a stimulant, depressant, or hallucinogenic effect on the
10		central nervous system that is substantially similar to or greater than the
11		stimulant, depressant, or hallucinogenic effect on the central nervous
12		system of a controlled substance in Schedule I or II.
13	(b) Such	n term does not include:
14	1.	Any substance for which there is an approved new drug application;
15	2.	With respect to a particular person, any substance if an exemption is in
16		effect for investigational use for that person pursuant to federal law to
17		the extent conduct with respect to such substance is pursuant to such
18		exemption; or
19	3.	Any substance to the extent not intended for human consumption before
20		the exemption described in subparagraph 2. of this paragraph takes
21		effect with respect to that substance;
22	(10) "Counterfo	eit substance" means a controlled substance which, or the container or
23	labeling o	f which, without authorization, bears the trademark, trade name, or other
24	identifying	g mark, imprint, number, or device, or any likeness thereof, of a
25	manufactu	arer, distributor, or dispenser other than the person who in fact
26	manufactu	ared, distributed, or dispensed the substance;

(11) "Dispense" means to deliver a controlled substance to an ultimate user or research

1		subje	ect by or pursuant to the lawful order of a practitioner, including the packaging,
2		label	ing, or compounding necessary to prepare the substance for that delivery;
3	(12)	"Dis	penser" means a person who lawfully dispenses a Schedule II, III, IV, or V
4		conti	rolled substance to or for the use of an ultimate user;
5	(13)	"Dis	tribute" means to deliver other than by administering or dispensing a controlled
6		subs	tance;
7	(14)	"Dos	sage unit" means a single pill, capsule, ampule, liquid, or other form of
8		admi	inistration available as a single unit;
9	(15)	"Dru	g" means:
10		(a)	Substances recognized as drugs in the official United States Pharmacopoeia,
11			official Homeopathic Pharmacopoeia of the United States, or official National
12			Formulary, or any supplement to any of them;
13		(b)	Substances intended for use in the diagnosis, care, mitigation, treatment, or
14			prevention of disease in man or animals;
15		(c)	Substances (other than food) intended to affect the structure or any function of
16			the body of man or animals; and
17		(d)	Substances intended for use as a component of any article specified in this
18			subsection.
19		It do	es not include devices or their components, parts, or accessories;
20	(16)	"Fen	tanyl" means a substance containing any quantity of fentanyl, or any of its
21		salts	, isomers, or salts of isomers, but does not include any quantity of fentanyl
22		<u>foun</u>	d on testing equipment as allowed in subsection (7) of Section 2 of this Act;
23	(17)	"Fen	tanyl derivative" means a substance containing any quantity of any chemical
24		comp	pound, except compounds specifically scheduled as controlled substances by
25		statu	te or by administrative regulation pursuant to this chapter, which is structurally
26		deriv	ved from 1-ethyl-4-(N-phenylamido) piperadine:
27		(a)	By substitution:

Page 3 of 19
XXXX 2/16/2023 1:58 PM Jacketed

1			1.	At the 2-position of the 1-ethyl group with a phenyl, furan, thiophene, or
2				ethyloxotetrazole ring system; and
3			2.	Of the terminal amido hydrogen atom with an alkyl, alkoxy, cycloalkyl,
4				or furanyl group; and
5		(b)	Whi	ich may be further modified in one (1) or more of the following ways:
6			1.	By substitution on the N-phenyl ring to any extent with alkyl, alkoxy,
7				haloalkyl, hydroxyl, or halide substituents;
8			2.	By substitution on the piperadine ring to any extent with alkyl, allyl,
9				alkoxy, hydroxy, or halide substituents at the 2-, 3-, 5-, and/or 6-
10				positions;
11			3.	By substitution on the piperadine ring to any extent with a phenyl,
12				alkoxy, or carboxylate ester substituent at the 4- position; or
13			4.	By substitution on the 1-ethyl group to any extent with alkyl, alkoxy, or
14				hydroxy substituents;
15	(18)	"Go	od fa	ith prior examination," as used in KRS Chapter 218A and for criminal
16		pros	ecutio	on only, means an in-person medical examination of the patient conducted
17		by t	he pi	rescribing practitioner or other health-care professional routinely relied
18		upor	n in t	the ordinary course of his or her practice, at which time the patient is
19		phys	sically	y examined and a medical history of the patient is obtained. "In-person"
20		inclu	ıdes t	elehealth examinations. This subsection shall not be applicable to hospice
21		prov	iders	licensed pursuant to KRS Chapter 216B;
22	(19)	"Haz	zardo	us chemical substance" includes any chemical substance used or intended
23		for u	ise in	the illegal manufacture of a controlled substance as defined in this section
24		or tl	he ill	egal manufacture of methamphetamine as defined in KRS 218A.1431,
25		whic	ch:	
26		(a)	Pose	es an explosion hazard;
27		(b)	Pose	es a fire hazard; or

Page 4 of 19 XXXX 2/16/2023 1:58 PM Jacketed

1 (c) Is poisonous or injurious if har	ndled, swallowed, or inhaled;
--	-------------------------------

- 2 (20) "Heroin" means a substance containing any quantity of heroin, or any of its salts,
- 3 isomers, or salts of isomers;
- 4 "Hydrocodone combination product" means a drug with: (21)
- Not more than three hundred (300) milligrams of dihydrocodeinone, or any of 5 (a) 6 its salts, per one hundred (100) milliliters or not more than fifteen (15) 7 milligrams per dosage unit, with a fourfold or greater quantity of an
- 8 isoquinoline alkaloid of opium; or
- 9 Not more than three hundred (300) milligrams of dihydrocodeinone, or any of 10 its salts, per one hundred (100) milliliters or not more than fifteen (15) 11 milligrams per dosage unit, with one (1) or more active, nonnarcotic
- 12 ingredients in recognized therapeutic amounts;
- (22) "Immediate precursor" means a substance which is the principal compound 13 14 commonly used or produced primarily for use, and which is an immediate chemical 15 intermediary used or likely to be used in the manufacture of a controlled substance 16 or methamphetamine, the control of which is necessary to prevent, curtail, or limit manufacture; 17
- 18 "Industrial hemp" has the same meaning as in KRS 260.850; (23)
- 19 "Industrial hemp products" has the same meaning as in KRS 260.850;
- 20 "Intent to manufacture" means any evidence which demonstrates a person's
- 21 conscious objective to manufacture a controlled substance or methamphetamine.
- 22 Such evidence includes but is not limited to statements and a chemical substance's
- 23 usage, quantity, manner of storage, or proximity to other chemical substances or
- 24 equipment used to manufacture a controlled substance or methamphetamine;
- 25 "Isomer" means the optical isomer, except the Cabinet for Health and Family
- 26 Services may include the optical, positional, or geometric isomer to classify any
- 27 substance pursuant to KRS 218A.020;

(27)	"Manufacture," except as provided in KRS 218A.1431, means the production,
	preparation, propagation, compounding, conversion, or processing of a controlled
	substance, either directly or indirectly by extraction from substances of natural
	origin or independently by means of chemical synthesis, or by a combination of
	extraction and chemical synthesis, and includes any packaging or repackaging of
	the substance or labeling or relabeling of its container except that this term does not
	include activities:

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

27

- By a practitioner as an incident to his or her administering or dispensing of a (a) controlled substance in the course of his or her professional practice;
- (b) By a practitioner, or by his or her authorized agent under his supervision, for the purpose of, or as an incident to, research, teaching, or chemical analysis and not for sale; or
- By a pharmacist as an incident to his or her dispensing of a controlled (c) substance in the course of his or her professional practice;
- (28) "Marijuana" means all parts of the plant Cannabis sp., whether growing or not; the seeds thereof; the resin extracted from any part of the plant; and every compound, manufacture, salt, derivative, mixture, or preparation of the plant, its seeds or resin or any compound, mixture, or preparation which contains any quantity of these substances. The term "marijuana" does not include:
  - Industrial hemp that is in the possession, custody, or control of a person who (a) holds a license issued by the Department of Agriculture permitting that person to cultivate, handle, or process industrial hemp;
  - Industrial hemp products that do not include any living plants, viable seeds, (b) leaf materials, or floral materials;
- (c) The substance cannabidiol, when transferred, dispensed, or administered 26 pursuant to the written order of a physician practicing at a hospital or associated clinic affiliated with a Kentucky public university having a college

1			or school of medicine;			
2		(d)	For persons participating in a clinical trial or in an expanded access program,			
3			a drug or substance approved for the use of those participants by the United			
4			States Food and Drug Administration;			
5		(e)	A cannabidiol product derived from industrial hemp, as defined in KRS			
6			260.850;			
7		(f)	For the purpose of conducting scientific research, a cannabinoid product			
8			derived from industrial hemp, as defined in KRS 260.850; or			
9		(g)	A cannabinoid product approved as a prescription medication by the United			
10			States Food and Drug Administration;			
11	(29)	"Me	dical history," as used in KRS Chapter 218A and for criminal prosecution only,			
12		mea	ns an accounting of a patient's medical background, including but not limited to			
13		prio	r medical conditions, prescriptions, and family background;			
14	(30)	"Me	dical order," as used in KRS Chapter 218A and for criminal prosecution only,			
15		mea	ns a lawful order of a specifically identified practitioner for a specifically			
16		iden	identified patient for the patient's health-care needs. "Medical order" may or may			
17		not i	nclude a prescription drug order;			
18	(31)	"Me	dical record," as used in KRS Chapter 218A and for criminal prosecution only,			
19		mea	ns a record, other than for financial or billing purposes, relating to a patient,			
20		kept	by a practitioner as a result of the practitioner-patient relationship;			
21	(32)	"Me	thamphetamine" means any substance that contains any quantity of			
22		meth	namphetamine, or any of its salts, isomers, or salts of isomers;			
23	(33)	"Naı	rcotic drug" means any of the following, whether produced directly or indirectly			
24		by e	extraction from substances of vegetable origin, or independently by means of			
25		chen	nical synthesis, or by a combination of extraction and chemical synthesis:			
26		(a)	Opium and opiate, and any salt, compound, derivative, or preparation of			

opium or opiate;

27

(b)	Any salt, compound, isomer, derivative, or preparation thereof which is
	chemically equivalent or identical with any of the substances referred to in
	paragraph (a) of this subsection, but not including the isoquinoline alkaloids
	of opium;

(c) Opium poppy and poppy straw;

1

2

3

4

5

6

7

8

- (d) Coca leaves, except coca leaves and extracts of coca leaves from which cocaine, ecgonine, and derivatives of ecgonine or their salts have been removed;
- 9 (e) Cocaine, its salts, optical and geometric isomers, and salts of isomers;
- 10 (f) Ecgonine, its derivatives, their salts, isomers, and salts of isomers; and
- 11 (g) Any compound, mixture, or preparation which contains any quantity of any of 12 the substances referred to in paragraphs (a) to (f) of this subsection;
- 13 (34) "Opiate" means any substance having an addiction-forming or addiction-sustaining
  14 liability similar to morphine or being capable of conversion into a drug having
  15 addiction-forming or addiction-sustaining liability. It does not include, unless
  16 specifically designated as controlled under KRS 218A.020, the dextrorotatory
  17 isomer of 3-methoxy-n-methylmorphinan and its salts (dextromethorphan). It does
  18 include its racemic and levorotatory forms;
- 19 (35) "Opium poppy" means the plant of the species papaver somniferum L., except its 20 seeds;
- 21 (36) "Person" means individual, corporation, government or governmental subdivision 22 or agency, business trust, estate, trust, partnership or association, or any other legal 23 entity;
- 24 (37) "Physical injury" has the same meaning it has in KRS 500.080;
- 25 (38) "Poppy straw" means all parts, except the seeds, of the opium poppy, after mowing;
- 26 (39) "Pharmacist" means a natural person licensed by this state to engage in the practice 27 of the profession of pharmacy;

(40) "Practitioner" means a physician, dentist, podiatrist, veterinarian, scientific investigator, optometrist as authorized in KRS 320.240, advanced practice registered nurse as authorized under KRS 314.011, physician assistant as authorized under KRS 311.858, or other person licensed, registered, or otherwise permitted by state or federal law to acquire, distribute, dispense, conduct research with respect to, or to administer a controlled substance in the course of professional practice or research in this state. "Practitioner" also includes a physician, dentist, podiatrist, veterinarian, or advanced practice registered nurse authorized under KRS 314.011 who is a resident of and actively practicing in a state other than Kentucky and who is licensed and has prescriptive authority for controlled substances under the professional licensing laws of another state, unless the person's Kentucky license has been revoked, suspended, restricted, or probated, in which case the terms of the Kentucky license shall prevail;

- (41) "Practitioner-patient relationship," as used in KRS Chapter 218A and for criminal prosecution only, means a medical relationship that exists between a patient and a practitioner or the practitioner's designee, after the practitioner or his or her designee has conducted at least one (1) good faith prior examination;
- 18 (42) "Prescription" means a written, electronic, or oral order for a drug or medicine, or
  19 combination or mixture of drugs or medicines, or proprietary preparation, signed or
  20 given or authorized by a medical, dental, chiropody, veterinarian, optometric
  21 practitioner, or advanced practice registered nurse, and intended for use in the
  22 diagnosis, cure, mitigation, treatment, or prevention of disease in man or other
  23 animals;
- 24 (43) "Prescription blank," with reference to a controlled substance, means a document that meets the requirements of KRS 218A.204 and 217.216;
- 26 (44) "Presumptive probation" means a sentence of probation not to exceed the maximum 27 term specified for the offense, subject to conditions otherwise authorized by law,

that is presumed to be the appropriate sentence for certain offenses designated in this chapter, notwithstanding contrary provisions of KRS Chapter 533. That presumption shall only be overcome by a finding on the record by the sentencing court of substantial and compelling reasons why the defendant cannot be safely and effectively supervised in the community, is not amenable to community-based treatment, or poses a significant risk to public safety;

- (45) "Production" includes the manufacture, planting, cultivation, growing, or harvesting of a controlled substance;
- (46) "Recovery program" means an evidence-based, nonclinical service that assists individuals and families working toward sustained recovery from substance use and other criminal risk factors. This can be done through an array of support programs and services that are delivered through residential and nonresidential means;
  - (47) "Salvia" means Salvia divinorum or Salvinorin A and includes all parts of the plant presently classified botanically as Salvia divinorum, whether growing or not, the seeds thereof, any extract from any part of that plant, and every compound, manufacture, derivative, mixture, or preparation of that plant, its seeds, or its extracts, including salts, isomers, and salts of isomers whenever the existence of such salts, isomers, and salts of isomers is possible within the specific chemical designation of that plant, its seeds, or extracts. The term shall not include any other species in the genus salvia;
- (48) "Second or subsequent offense" means that for the purposes of this chapter an offense is considered as a second or subsequent offense, if, prior to his or her conviction of the offense, the offender has at any time been convicted under this chapter, or under any statute of the United States, or of any state relating to substances classified as controlled substances or counterfeit substances, except that a prior conviction for a nontrafficking offense shall be treated as a prior offense only when the subsequent offense is a nontrafficking offense. For the purposes of

this section, a conviction voided under KRS 218A.275 or 218A.276 shall not constitute a conviction under this chapter;

- 3 (49) "Sell" means to dispose of a controlled substance to another person for consideration or in furtherance of commercial distribution;
- 5 (50) "Serious physical injury" has the same meaning it has in KRS 500.080;

13

14

15

16

17

18

19

20

21

22

23

24

25

26

27

- 6 "Synthetic cannabinoids or piperazines" means any chemical compound which is 7 not approved by the United States Food and Drug Administration or, if approved, 8 which is not dispensed or possessed in accordance with state and federal law, that 9 contains Benzylpiperazine (BZP); Trifluoromethylphenylpiperazine (TFMPP); 1,1-10 Dimethylheptyl-11-hydroxytetrahydrocannabinol (HU-210); 1-Butyl-3-(1-11 naphthoyl)indole; 1-Pentyl-3-(1-naphthoyl)indole; dexanabinol (HU-211); or any 12 compound in the following structural classes:
  - (a) Naphthoylindoles: Any compound containing a 3-(1-naphthoyl)indole structure with substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group, whether or not further substituted in the indole ring to any extent and whether or not substituted in the naphthyl ring to any extent. Examples of this structural class include but are not limited to JWH-015, JWH-018, JWH-019, JWH-073, JWH-081, JWH-122, JWH-200, and AM-2201;
  - (b) Phenylacetylindoles: Any compound containing a 3-phenylacetylindole structure with substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group whether or not further substituted in the indole ring to any extent and whether or not substituted in the phenyl ring to any extent. Examples of this structural class include but are not limited to JWH-167, JWH-250, JWH-251, and RCS-8;

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

26

27

(c)	(c) Benzoylindoles: Any compound containing a 3-(benzoyl)indole structure					
	substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl,					
	alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-					
	piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group whether or not further					
	substituted in the indole ring to any extent and whether or not substituted in					
	the phenyl ring to any extent. Examples of this structural class include but are					
	not limited to AM-630, AM-2233, AM-694, Pravadoline (WIN 48,098), and					
	RCS-4;					

- (d) Cyclohexylphenols: compound containing 2-(3-Any a hydroxycyclohexyl)phenol structure with substitution at the 5-position of the ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group whether or not substituted in the cyclohexyl ring to any extent. Examples of this structural class include but are not limited to CP 47,497 and its C8 homologue (cannabicyclohexanol);
- (e) Naphthylmethylindoles: Any compound containing a 1H-indol-3-yl-(1-naphthyl)methane structure with substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group whether or not further substituted in the indole ring to any extent and whether or not substituted in the naphthyl ring to any extent. Examples of this structural class include but are not limited to JWH-175, JWH-184, and JWH-185;
- (f) Naphthoylpyrroles: Any compound containing a 3-(1-naphthoyl)pyrrole structure with substitution at the nitrogen atom of the pyrrole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group whether or not further substituted in the pyrrole ring to any extent and whether or not substituted in

the naphthyl ring to any extent. Examples of this structural class include but are not limited to JWH-030, JWH-145, JWH-146, JWH-307, and JWH-368;

- (g) Naphthylmethylindenes: Any compound containing a 1-(1-naphthylmethyl)indene structure with substitution at the 3-position of the indene ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group whether or not further substituted in the indene ring to any extent and whether or not substituted in the naphthyl ring to any extent. Examples of this structural class include but are not limited to JWH-176;
- (h) Tetramethylcyclopropanoylindoles: Any compound containing a 3-(1-tetramethylcyclopropoyl)indole structure with substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group, whether or not further substituted in the indole ring to any extent and whether or not further substituted in the tetramethylcyclopropyl ring to any extent. Examples of this structural class include but are not limited to UR-144 and XLR-11;
- (i) Adamantoylindoles: Any compound containing a 3-(1-adamantoyl)indole structure with substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, 1-(N-methyl-2-piperidinyl)methyl, or 2-(4-morpholinyl)ethyl group, whether or not further substituted in the indole ring to any extent and whether or not substituted in the adamantyl ring system to any extent. Examples of this structural class include but are not limited to AB-001 and AM-1248; or
- (j) Any other synthetic cannabinoid or piperazine which is not approved by the United States Food and Drug Administration or, if approved, which is not dispensed or possessed in accordance with state and federal law;

I	(52) "Synthetic cath	ninones" means any chemical compound which is not approved by
2	the United Sta	ites Food and Drug Administration or, if approved, which is not
3	dispensed or p	possessed in accordance with state and federal law (not including
4	bupropion or co	ompounds listed under a different schedule) structurally derived from
5	2-aminopropan	1-1-one by substitution at the 1-position with either phenyl, naphthyl,
6	or thiophene ri	ng systems, whether or not the compound is further modified in one
7	(1) or more of	the following ways:

1

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

- By substitution in the ring system to any extent with alkyl, alkylenedioxy, (a) alkoxy, haloalkyl, hydroxyl, or halide substituents, whether or not further substituted in the ring system by one (1) or more other univalent substituents. Examples of this class include but are not limited Methylenedioxycathinone (bk-MDA);
- By substitution at the 3-position with an acyclic alkyl substituent. Examples (b) of this class include but are not limited to 2-methylamino-1-phenylbutan-1one (buphedrone);
- (c) By substitution at the 2-amino nitrogen atom with alkyl, dialkyl, benzyl, or methoxybenzyl groups, or by inclusion of the 2-amino nitrogen atom in a cyclic structure. Examples of this class include but are not limited to Dimethylcathinone, Ethcathinone, and  $\alpha$ -Pyrrolidinopropiophenone ( $\alpha$ -PPP); or
- (d) Any other synthetic cathinone which is not approved by the United States Food and Drug Administration or, if approved, is not dispensed or possessed in accordance with state or federal law;
- 24 (53) "Synthetic drugs" means any synthetic cannabinoids or piperazines or any synthetic 25 cathinones;
- 26 (54) "Telehealth" has the same meaning it has in KRS 311.550;
- 27 (55) "Tetrahydrocannabinols" means synthetic equivalents of the substances contained

1	in the plant, or in the resinous extractives of the plant Cannabis, sp. or synthetic
2	substances, derivatives, and their isomers with similar chemical structure and
3	pharmacological activity such as the following:
4	(a) Delta 1 cis or trans tetrahydrocannabinol, and their optical isomers:

- Delta 1 cis or trans tetrahydrocannabinol, and their optical isomers; (a)
- Delta 6 cis or trans tetrahydrocannabinol, and their optical isomers; and 5 (b)
- 6 Delta 3, 4 cis or trans tetrahydrocannabinol, and its optical isomers;
- 7 (56) "Traffic," except as provided in KRS 218A.1431, means to manufacture, distribute,
- 8 dispense, sell, transfer, or possess with intent to manufacture, distribute, dispense,
- 9 or sell a controlled substance;
- 10 "Transfer" means to dispose of a controlled substance to another person without
- 11 consideration and not in furtherance of commercial distribution; and
- 12 (58) "Ultimate user" means a person who lawfully possesses a controlled substance for
- 13 his or her own use or for the use of a member of his or her household or for
- 14 administering to an animal owned by him or her or by a member of his or her
- 15 household.
- 16 → Section 2. KRS 218A.500 is amended to read as follows:
- As used in this section and KRS 218A.510: 17
- 18 "Drug paraphernalia" means all equipment, products and materials of any kind (1)
- 19 which are used, intended for use, or designed for use in planting, propagating,
- 20 cultivating, growing, harvesting, manufacturing, compounding, converting,
- 21 producing, processing, preparing, testing, analyzing, packaging, repackaging,
- 22 storing, containing, concealing, injecting, ingesting, inhaling, or otherwise
- 23 introducing into the human body a controlled substance in violation of this chapter.
- 24 It includes but is not limited to:
- 25 Kits used, intended for use, or designed for use in planting, propagating,
- 26 cultivating, growing, or harvesting of any species of plant which is a
- 27 controlled substance or from which a controlled substance can be derived;

1	(b)	Kits used, intended for use, or designed for use in manufacturing,
2		compounding, converting, producing, processing, or preparing controlled
3		substances;
4	(c)	Isomerization devices used, intended for use, or designed for use in increasing
5		the potency of any species of plant which is a controlled substance;
6	(d)	Except as provided for in subsection (7), testing equipment used, intended for
7		use, or designed for use in [identifying, or in ]analyzing the strength,
8		effectiveness, or purity of controlled substances;
9	(e)	Scales and balances used, intended for use, or designed for use in weighing or
10		measuring controlled substances;
11	(f)	Diluents and adulterants, such as quinine hydrochloride, mannitol, mannite,
12		dextrose and lactose, used, intended for use, or designed for use in cutting
13		controlled substances;
14	(g)	Separation gins and sifters used, intended for use, or designed for use in
15		removing twigs and seeds from, or in otherwise cleaning or refining
16		marijuana;
17	(h)	Blenders, bowls, containers, spoons, and mixing devices used, intended for
18		use, or designed for use in compounding controlled substances;
19	(i)	Capsules, balloons, envelopes, and other containers used, intended for use, or
20		designed for use in packaging small quantities of controlled substances;
21	(j)	Containers and other objects used, intended for use, or designed for use in
22		storing or concealing controlled substances;
23	(k)	Hypodermic syringes, needles, and other objects used, intended for use, or
24		designed for use in parenterally injecting controlled substances into the human
25		body; and
26	(1)	Objects used, intended for use, or designed for use in ingesting, inhaling, or

Page 16 of 19
XXXX 2/16/2023 1:58 PM Jacketed

otherwise introducing marijuana, cocaine, hashish, or hashish oil into the

27

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

17

18

19

20

21

22

23

24

25

human body, such as: metal, wooden, acrylic, glass, stone, plastic, or ceramic pipes with or without screens, permanent screens, hashish heads, or punctured metal bowls; water pipes; carburetion tubes and devices; smoking and carburetion masks; roach clips which mean objects used to hold burning material, such as marijuana cigarettes, that have become too small or too short to be held in the hand; miniature cocaine spoons, and cocaine vials; chamber pipes; carburetor pipes; electric pipes; air-driven pipes; chillums; bongs; ice pipes or chillers.

- It is unlawful for any person to use, or to possess with intent to use, drug paraphernalia for the purpose of planting, propagating, cultivating, growing, harvesting, manufacturing, compounding, converting, producing, processing, preparing, testing, analyzing, packing, repacking, storing, containing, concealing, injecting, ingesting, inhaling, or otherwise introducing into the human body a controlled substance in violation of this chapter.
- It is unlawful for any person to deliver, possess with intent to deliver, or (3) 16 manufacture with intent to deliver, drug paraphernalia, knowing, or under circumstances where one reasonably should know, that it will be used to plant, propagate, cultivate, grow, harvest, manufacture, compound, convert, produce, process, prepare, test, analyze, pack, repack, store, contain, conceal, inject, ingest, inhale, or otherwise introduce into the human body a controlled substance in violation of this chapter.
  - (4) It is unlawful for any person to place in any newspaper, magazine, handbill, or other publication any advertisement, knowing, or under circumstances where one reasonably should know, that the purpose of the advertisement, in whole or in part, is to promote the sale of objects designed or intended for use as drug paraphernalia.
- 26 (5) (a) This section shall not prohibit a local health department from operating a 27 substance abuse treatment outreach program which allows participants to

1	exchange	hypodermic	needles	and	svringes
1	CACHAIIge	ny podernine	necures	ana	symmets.

2

3

4

5

6

7

8

9

10

21

22

23

- (b) To operate a substance abuse treatment outreach program under this subsection, the local health department shall have the consent, which may be revoked at any time, of the local board of health and:
  - The legislative body of the first or home rule class city in which the program would operate if located in such a city; and
  - 2. The legislative body of the county, urban-county government, or consolidated local government in which the program would operate.
  - (c) Items exchanged at the program shall not be deemed drug paraphernalia under this section while located at the program.
- 11 (6) (a) Prior to searching a person, a person's premises, or a person's vehicle, a peace 12 officer may inquire as to the presence of needles or other sharp objects in the 13 areas to be searched that may cut or puncture the officer and offer to not 14 charge a person with possession of drug paraphernalia if the person declares 15 to the officer the presence of the needle or other sharp object. If, in response 16 to the offer, the person admits to the presence of the needle or other sharp 17 object prior to the search, the person shall not be charged with or prosecuted 18 for possession of drug paraphernalia for the needle or sharp object or for 19 possession of a controlled substance for residual or trace drug amounts 20 present on the needle or sharp object.
  - (b) The exemption under this subsection shall not apply to any other drug paraphernalia that may be present and found during the search or to controlled substances present in other than residual or trace amounts.
- 24 (7) (a) This section shall not prohibit the retail sale of hypodermic syringes and needles without a prescription in pharmacies.
- 26 (b) Hypodermic syringe and needle inventory of a pharmacy shall not be deemed 27 drug paraphernalia under this section.

1	(c) Testing equipment utilized in determining whether a controlled substance
2	contains chemicals, toxic substances, or hazardous compounds which can
3	cause physical harm or death shall not be deemed drug paraphernalia
4	under this section. This includes but is not limited to fentanyl test strips.
5	(8) Any person who violates any provision of this section shall be guilty of a Class A
6	misdemeanor.
7	→ Section 3. The Cabinet for Health and Family Services in coordination with the
8	Justice and Public Safety Cabinet shall conduct or have conducted a Fentanyl Education
9	and Awareness campaign. The campaign shall begin no later than 90 days after the
10	effective date of this Act.